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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/202,054	12/07/1998	AUDREY GODDARD	P1154R2	2403
7590	10/26/2004		EXAMINER	
GINGER R DREGER			SPECTOR, LORRAINE	
GENENTECH INC				
1 DNA WAY			ART UNIT	PAPER NUMBER
SOUTH SAN FRANCISCO, CA 940804990			1647	

DATE MAILED: 10/26/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary	Application No.	Applicant(s)
	09/202,054	GODDARD ET AL.
	Examiner	Art Unit
	Lorraine Spector, Ph.D.	1647

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --
Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM
 THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
 - If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
 - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
 - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) Responsive to communication(s) filed on 16 August 2004.
- 2a) This action is **FINAL**. 2b) This action is non-final.
- 3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) Claim(s) 28-30,48-50 and 54-57 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) Claim(s) _____ is/are allowed.
- 6) Claim(s) 28-30 and 48-50 and 54-57 is/are rejected.
- 7) Claim(s) _____ is/are objected to.
- 8) Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) The specification is objected to by the Examiner.
- 10) The drawing(s) filed on _____ is/are: a) accepted or b) objected to by the Examiner.
 Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
 Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
 a) All b) Some * c) None of:
 1. Certified copies of the priority documents have been received.
 2. Certified copies of the priority documents have been received in Application No. _____.
 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|--|---|
| 1) <input type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413) |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | Paper No(s)/Mail Date. _____ |
| 3) <input type="checkbox"/> Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)
Paper No(s)/Mail Date _____ | 5) <input type="checkbox"/> Notice of Informal Patent Application (PTO-152) |
| | 6) <input type="checkbox"/> Other: _____ |

Continued Examination Under 37 CFR 1.114

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 12/9/2003 has been entered.

Claims 28-30 and 48-50 and 54 are pending and under consideration.

Applicants are requested to provide sufficient identifying information regarding Exhibit "C" submitted with the response of 12/9/2003 so that the Examiner can properly make that reference of record.

The Bazan declaration filed 12/9/2003 under 37 CFR 1.132 is insufficient to overcome the rejection of claims 28-30, 48-50 and 54 35 U.S.C. §101 and §112 as set forth in the last Office action because:

Dr. Bazan refers to the specification at Figure 7B and page 7, lines 8-23 as showing that PRO285 has significant homology to the IL-1 receptor domain that is necessary for signaling via NF- κ B and that such would be predictive of NF- κ B-mediated signaling by PRO285. This argument has been fully considered but is not deemed persuasive because Figure 7 shows an alignment of IL-1R with TLR2 in the region critical for IL-1R signaling. However, examination of Figure 7B reveals that six residues are indicated as being "essential for IL-1R signaling", and only three of those six are conserved in TLR2, which had not, as of the filing date of this application, been shown to signal via NF- κ B. The declaration provides, for the first time, an alignment of the analogous region of PRO285 with both IL-1R and TLR2. The Examiner notes that of the six essential residues for IL-1R signaling, only *two* are conserved in PRO285. This amount of conservation is not persuasive of conservation of function. Therefore, based upon the information in the specification as originally filed, it does not appear that it would lead a person

of ordinary skill in the art to the conclusion that PRO285 would be expected to have NF- $\kappa\beta$ signaling activity.

At paragraphs 6-7, Dr. Bazan refers to an article by Rock et al., PNAS 95:588-593, previously of record, as teaching that it would be expected that PRO285 would signal via NF- $\kappa\beta$. This argument has been fully considered but is not deemed persuasive because while the Rock article clearly shows that five TLRs were evolutionarily related to the *Drosophila* Toll receptors, there is no clear indication that they would be expected to activate NF- $\kappa\beta$, nor is there disclosure or discussion of PRO285. Declarant is arguing based on two steps removed from the data; the argument is that the function of IL-1R was known, that TLRs are similar to IL-1R and thus might share function, and that PRO285 is similar to TLRs. Further, mere signaling via NF- $\kappa\beta$ would not be indicative of utility, for reasons cited below. While the Rock et al. paper states that the TLRs “could constitute an important and unrecognized component of innate immunity in humans”, it also states, in the very next sentence, that “Intriguingly, the evolutionary retention of TLRs in vertebrates may indicate another role- akin to Toll in the dorsoventralization of the *Drosophila* embryo- as regulators of early morphogenetic patterning. Finally, the Rock paper does not disclose or discuss PRO285. Thus, based upon the Rock paper, it would seem that it was *not* predictable that PRO285 was involved in innate immunity via signaling via NF- $\kappa\beta$, and that there were at least two very distinct ideas as to what the receptors disclosed therein might do. Finally, the fact that TLR4 was confirmed as signaling via NF- $\kappa\beta$ does not have bearing on PRO285.

At paragraph 8, Dr. Bazan concludes that the person of ordinary skill in the art could, reading the specification, reasonably conclude that PRO285 signals via NF- $\kappa\beta$, and that “antibodies to PRO285 could be made and used in accordance with routine techniques to modulate such activity.” This argument has been fully considered but is not deemed persuasive because it is the opinion of the declarant, and is not supported by the facts and evidence of record, specifically: (a) As stated above, as essential residues of the IL-1R signaling domain are *not* conserved in PRO285, the Examiner does not agree with the assertion that the person of ordinary skill in the art, reading the specification as originally filed, would find it predictable that PRO285 signals via NF- $\kappa\beta$. (b) The Rock publication indicates that TLRs *might* signal via NF-

$\kappa\beta$, or alternatively, *might* be involved in early morphogenic patterning, and does not disclose or discuss PRO285. (c) Even *if* it were predictable that PRO285 signaled via NF- $\kappa\beta$, such is not a utility, in and of itself. NF- $\kappa\beta$ is an intracellular signaling molecule. Without knowing any ligand for PRO285 nor under what physiological circumstances the ligand binds to the receptor, the person of ordinary skill in the art would not know how to use the receptor for its NF- $\kappa\beta$ signaling activity, that is, under what circumstances it would be desirable to stimulate or inhibit PRO285.

Claim Rejections - 35 USC § 101 and §112, first paragraph

35 U.S.C. 101 reads as follows:

Whoever invents or discovers any new and useful process, machine, manufacture, or composition of matter, or any new and useful improvement thereof, may obtain a patent therefor, subject to the conditions and requirements of this title.

Claims 28-30, 48-50 and 54 remain rejected under 35 U.S.C. 101 because the claimed invention is not supported by either a specific, substantial and credible asserted utility or a well established utility, for reasons of record in the previous Office Action mailed 9/16/2002 at pages 2-4. Applicants traversal of this rejection in the paper submitted 12/9/2003 have been fully considered but are not deemed persuasive.

At page 2, applicants argue that the Examiner is mis-applying the utility guidelines as published by the USPTO in the Federal Register, 60 FR 36263-02, published in 1995. The Examiner points applicants to 66 FR 1092, published 1/5/2001, which sets for the standard not as specific and credible, but specific, *substantial* and credible. There is no *substantial* assertion of utility in this case, without which credibility cannot be assessed.

At page 3 of the response, applicants argue that the specification teaches that PRO285 polypeptide signaling activates NF- $\kappa\beta$ and that antibodies to PRO285 can be used to modulate this activity. This argument has been fully considered but is not deemed persuasive because without knowing under what conditions PRO285 so signals, the use of such antibodies would merely constitute further research to determine the function and biological role of PRO285,

which is not considered to constitute a utility under 35 U.S.C. §101. Applicants have submitted with their response a paper by Jurk et al., (Nature Immunology 3:499, 2002), which demonstrates that even four years after the filing date of the instant application, the biological function of PRO285, by applicants assertion later designated TLR7, was unknown. Specifically, Jurk et al. teach that different TLRs signal in response to different stimuli; TLR2, 4 and 5 in response to peptidoglycan, lipopolysaccharide and flagellin, respectively, TLR6 in conjunction with TLR2 in response to lipoproteins from mycoplasma, TLR9 in response to bacterial DNA containing unmethylated CpG motifs, and TLR3 in response to dsRNA (see abstract). The Jurk paper goes on to state that “The natural ligands for TLR1, TLR7, TLR8 and TLR10 are not known, although a synthetic compound with antiviral activity has not been described as a ligand for TLR7. Thus, even four years after the filing date of the instant application, the role of TLR7, aka PRO285, was unknown, and the receptor was merely a subject for further research. This paper supports the Examiner’s position that even *if* one were to accept, on the basis of the specification as originally filed, that PRO285 signals via NF- $\kappa\beta$, that such does not confer any specific, substantial and credible utility upon the protein, nor upon antibodies that bind to it.

Applicants repeatedly state in their argument that antibodies to PRO285 can be used to “modulate” NF- $\kappa\beta$ signaling, however, the specification fails to teach how to do so, the claims require not such activity, and it is not clear what utility “modulation” of such signaling has if one has no idea in response to what stimulus such signaling occurs in nature.

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claims 28-30, 48-50 and 54 also remain and claims 55-57 are rejected under 35 U.S.C. 112, first paragraph. Specifically, since the claimed invention is not supported by either a specific, substantial and credible asserted utility or a well established utility for the reasons set forth above, one skilled in the art clearly would not know how to use the claimed invention.

Applicants arguments of this rejection have been fully addressed above.

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

Claims 28 and 48 remain rejected under 35 U.S.C. 102(b) as being anticipated by Ruggeri et al., WO 91/09614.

Ruggeri et al. disclose a 19 residue peptide that matches SEQ ID NO: 2 at positions 704-712, a 9/15 match; see the third peptide listed in claim 1. At page 19 and in claim 65, antibodies to such peptides are disclosed and claimed.

Applicants argue that the three dimensional antigenic determinant of the 19 residue platelet membrane glycoprotein Ib peptide is not necessarily present in PRO285. This argument has been fully considered but is not deemed persuasive because according to "Exhibit C" submitted by applicants, although antibodies raised against fragments generally have higher affinities for the fragment to which they were raised than to the native protein, (page 247)they nonetheless "show extensive cross-reactions with native proteins" (page 248). Further, at page 249, the reference goes on to state that "antipeptide antibodies have proved to be very powerful reagents", and can be used to immunoprecipitate previously unisolated native proteins, to isolate previously unidentified gene products of new genes, and in detecting post-translational processing, are useful in probing structure-function relationships, and can be used to block protein binding. Accordingly, the reference provided by applicants teaches that one would reasonably expect an antibody raised against Ruggeri's peptide to bind to PRO285. Since the Office does not have the facilities for examining and comparing applicants' antibodies with those of the prior art, the burden is on applicant to show a novel or unobvious difference between the claimed product and the product of the prior art (i.e., that the protein of the prior art does not possess the same material structural and functional characteristics of the claimed protein). See In

Art Unit: 1647

re Best, 562 F.2d 1252, 195 USPQ 430 (CCPA 1977) and Ex parte Gray, 10 USPQ 2d 1922 1923 (PTO Bd. Pat. App. & Int.).

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

Claims 29 and 49 remain rejected under 35 U.S.C. 103(a) as being unpatentable over Ruggeri et al., WO 91/09614, in view of Coughlin, U.S. Patent Number 5,256,766 for reasons of record. Applicants argument of Ruggeri is found not persuasive, for reasons cited above. Applicants have presented no separate argument of the obviousness rejection.

Claims 50 and 54 remain rejected under 35 U.S.C. 103(a) as being unpatentable over Ruggeri et al., WO 91/09614, in view of Coughlin, U.S. Patent Number 5,256,766, and further in view of U.S. Patent Number 4,946,778 (Ladner et al.) for reasons of record. Applicants argument of Ruggeri is found not persuasive, for reasons cited above. Applicants have presented no separate argument of the obviousness rejection.

Conclusion

No claim is allowed.

All claims are drawn to the same invention claimed in the application prior to the entry of the submission under 37 CFR 1.114 and could have been finally rejected on the grounds and art of record in the next Office action if they had been entered in the application prior to entry under 37 CFR 1.114. Accordingly, **THIS ACTION IS MADE FINAL** even though it is a first action after the filing of a request for continued examination and the submission under 37 CFR 1.114. See MPEP § 706.07(b). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

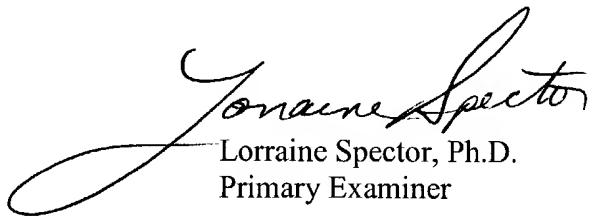
Any inquiry concerning this communication or earlier communications from the Examiner should be directed to Lorraine M. Spector. Dr. Spector can normally be reached Monday through Friday, 9:00 A.M. to 5:30 P.M. ***Effective 1/21/2004, Dr. Spector's telephone number is 571-272-0893.***

If attempts to reach the Examiner by telephone are unsuccessful, the Examiner's supervisor, Dr. Gary L. Kunz. ***Effective 1/21/2004, Dr. Kunz' telephone number is 571-272-0887.***

Certain papers related to this application may be submitted to Group 1800 by facsimile transmission. The faxing of such papers must conform with the notices published in the Official Gazette, 1156 OG 61 (November 16, 1993) and 1157 OG 94 (December 28, 1993) (see 37 C.F.R. § 1.6(d)). NOTE: If Applicant does submit a paper by fax, the original signed copy should be retained by applicant or applicant's representative. NO DUPLICATE COPIES SHOULD BE SUBMITTED so as to avoid the processing of duplicate papers in the Office.

Official papers filed by fax should be directed to (703) 872-9306 (before final rejection) or (703)872-9307 (after final). Faxed draft or informal communications with the examiner should be directed to **571-273-0893**.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).



Lorraine Spector
Lorraine Spector, Ph.D.
Primary Examiner